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AMENDMENTS TO THE CLAIMS

The following Listing of Claims replaces all prior versions, and listings, of claims in this Application.

LISTING OF CLAIMS

1-3 (cancelled)

- (currently amended) A method of promoting the treatment of intestinal cancer damage therapy in a patient comprising administering to the patient an effective amount of a pharmaceutically acceptable composition comprising (a) a non-naturally occurring polypeptide having an amino acid sequence according to the formula X1 H X2 D G S F S D E MNTX3LDX4LAX5X6DFINWLX7X8TKITDX9 His Xaa2 D Asp Gly Ser Phe Ser Asp Glu Met Asn Thr Xaa3 Leu Asp Xaa4 Leu Ala Xaa5 Xaa⁶ D Asp Phe Ile Asn Trp Leu Xaa⁷ Xaa⁸ Thr Lys Ile Thr Asp Xaa⁹ Xaa¹⁰ (SEQ ID NO:1), wherein Xaa2 is Ala or Gly; Xaa3 is Ile or Val; Xaa4 is Asn. Ser or His; Xaa⁵ is Ala or Thr; Xaa⁶ is Arg or Lys; Xaa⁷ is Ile or Leu: Xaa⁸ is Gln or His; Xaa9 is OH, Lys, or Arg, and Xaa10 is Lys, Arg, or is missing and optionally further including an N-terminal positioned sequence selected from Asp Phe Pro Glu Glu Val Ala Ile Val Glu Glu Leu Gly Arg Arg (SEQ ID NO:2), Asp Phe Pro Glu Glu Val Thr Ile Val Glu Glu Leu Gly Arg Arg (SEO ID NO:3), Asp Phe Pro Glu Glu Val Asn Ile Val Glu Glu Leu Arg Arg Arg (SEO ID NO:4), or a fragment of any of SEO ID NOS:2-4, and
- (b) a pharmaceutically acceptable combination of (i) an isotonic agent, (ii) a buffer, and (iii) a preservative, a surfactant, or a combination of a surfactant and a preservative, wherein (I) X1 is NH2, DFPEEVAIVEELGRR (SEQ ID NO:2), DFPEEVTIVEELGRR (SEQ ID NO:3), DFPEEVNIVEELRRR (SEQ ID NO:4), or a fragment of any of SEQ ID NOS:2-4; X2 is Ala-or-Gly; X3 is Ile-or-Val; X4-is Asn, Ser, or His; X5 is Ala-or-Thr; X6 is Arg or Lys; X7 is Ile-or-Leu; X8 is Gln or His; and X9 is OH, Lys, Arg, Arg Lys, Lys-Arg, Arg-Arg, or Lys-Lys-and (II) the solubility

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of the peptide, stability of the peptide, or both is significantly greater than the solubility and/or stability of the peptide without the combination to the patient.

(currently amended) The method of claim 4, wherein Xaa2 5. X2 represents Gly.

6-9 (cancelled)

(currently amended) A method of promoting the treatment of small bowel syndrome, Crohn's disease, ileitis, intestinal Inflammation, gastric ulceration, duodenal ulceration, inflammatory bowel-disease, or intestinal cancer damage therapy in a patient comprising administering an effective amount of the peptide of claim 6 a non-naturally occurring polypeptide having an amino acid sequence according to the formula His Xaa² D Asp Gly Ser Phe Ser Asp Glu Met Asn Thr Xaa³ Leu Asp Xaa⁴ Leu Ala Xaa⁵ Xaa⁶ D Asp Phe Ile Asn Trp Leu Xaa⁷ Xaa⁶ Thr Lys Ile Thr Asp Xaa⁹ Xaa10 (SEQ ID NO:1), wherein Xaa2 is Ala or Gly; Xaa3 is Ile or Val; Xaa4 is Asn. Ser or His; Xaa⁵ is Ala or Thr; Xaa⁶ is Arg or Lys; Xaa⁷ is Ile or Leu; Xaa⁸ is Gin or His; Xaa⁹ is OH, Lvs, or Arg, and Xaa¹⁰ is Lys, Arg, or is missing and optionally further including an N-terminal positioned sequence selected from Asp Phe Pro Glu Glu Val Ala Ile Val Glu Glu Leu Gly Arg Arg (SEQ ID NO:2), Asp Phe Pro Glu Glu Val Thr Ile Val Glu Glu Leu Gly Arg Arg (SEO ID NO:3), Asp Phe Pro Glu Glu Val Asn Ile Val Glu Glu Leu Arg Arg Arg (SEQ ID NO:4), or a fragment of any of SEQ ID NOS:2-4 X1 H-X2 D G-S-F S D E M N T X3 L D X4 L A X5 X6 D F I N W L X7 X8 T K I T D X9 (SEQ ID NO: 1), wherein (I) X1 is NH2, DFPEEVAIVEELGRR (SEQ ID NO:2), DFPEEVTIVEELGRR (SEQ-ID-NO:3), DFPEEVNIVEELRRR (SEQ-ID-NO:4), or a fragment of any of SEQ ID NOS:2 4; X2 is Ala or Gly; X3 is Ile or Val; X4 is Asn, Ser, or His; X5 is Ala or Thr; X6 is Arg or Lys; X7 is Ile or Leu; X8 is Gln or His; and X9 is OH; Lys, Arg, Arg-Lys, Lys Arg, Arg-Arg, or Lys-Lys to the patient.

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- The method of claim 10, wherein Xaa2 11. (currently amended) X2 represents Gly.
- The method of claim 10, wherein (previously presented) the composition comprises a preservative.
- The method of claim 12, wherein (previously presented) 13. the composition comprises a surfactant.
- The method of claim 4, wherein (previously presented) 14. the method comprises reconstituting a lyophilized composition comprising the non-naturally occurring polypeptide and preparing the pharmaceutically acceptable composition prior to administering the pharmaceutically acceptable composition to the patient.
- The method of claim 10, wherein (previously presented) 15. the method comprises reconstituting a lyophilized composition comprising the non-naturally occurring polypeptide prior to administering the polypeptide to the patient.